California Medical Device Recall Information

Recall Name
Zimmer Recalls **Zimmer M/L Taper with *Kinectiv* Technology Femoral Stems and Necks** **Due to Manufacturing Residues**

Recall Date	Product Description	Recalling Firm	Recall Reason
06/08/15	Zimmer M/L Taper with *Kinectiv* Technology Femoral Stems and Necks	**Zimmer, Inc.** Warsaw, IN	*Due to a process monitoring failure that led to higher than expected amounts of manufacturing residues left on the devices.*
Recall Class	**Product Identification**	**Distribution**	**Affected Dates**
I	List of Affected Lot Numbers and Product Descriptions	Nationwide	Manufacturing and distribution dates: March 31, 2015 through April 20, 2015

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm451936.htm